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INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

<p>(51) International Patent Classification : C07C 275/34, C07D 317/60, C07C 317/50, C07D 405/12, 407/12, 409/12, 417/12, 413/12, C07C 323/60, C07K 5/02, A61K 31/195, 31/36, 31/41</p>	<p>A3</p>	<p>(11) International Publication Number: WO 99/24398</p> <p>(43) International Publication Date: 20 May 1999 (20.05.99)</p>
<p>(21) International Application Number: PCT/GB98/03334</p> <p>(22) International Filing Date: 9 November 1998 (09.11.98)</p> <p>(30) Priority Data: 9723789.5 12 November 1997 (12.11.97) GB</p> <p>(71) Applicant (for all designated States except US): ZENECA LIMITED [GB/GB]; 15 Stanhope Gate, London W1Y 6LN (GB).</p> <p>(72) Inventors; and (75) Inventors/Applicants (for US only): BRITTAIN, David, Robert [GB/GB]; Alderley Park, Macclesfield, Cheshire SK10 4TG (GB). JOHNSTONE, Craig [GB/GB]; Alderley Park, Macclesfield, Cheshire SK10 4TG (GB).</p> <p>(74) Agent: BRYANT, Tracey; Zeneca Pharmaceuticals, Intellectual Property Dept., Mereside, Alderley Park, Macclesfield, Cheshire SK10 4TG (GB).</p>		<p>(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, HR, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).</p> <p>Published With international search report.</p> <p>(88) Date of publication of the international search report: 5 August 1999 (05.08.99)</p>
<p>(54) Title: UREA DERIVATIVES AND THEIR USE AS INTEGRIN INHIBITORS</p>		
<p style="text-align: right;">(II)</p>		
<p style="text-align: right;">(A)</p>		
<p style="text-align: right;">(B)</p>		
<p>(57) Abstract</p> <p>Compounds of formula (II) where R¹ is in the para or meta position and is (A); R² and R³ are each independently selected from hydrogen, nitro, C₁₋₆alkyl, C₃₋₆cycloalkyl, C₂₋₆alkenyl, C₂₋₆alkynyl, C₁₋₆alkoxy, C₁₋₆alkylamino, C₁₋₆dialkylamino, C₁₋₆alkyl(C₁₋₆alkoxy), C₁₋₆alkylaminoC₁₋₆alkyl, amino, cyano, halogeno, trifluoromethyl, -CO₂R¹² and -CONR¹²R¹³, where R¹² and R¹³ are independently selected from hydrogen or C₁₋₆alkyl, or R² and R³ together with the phenyl to which they are attached form a 9 or 10 membered bicyclic ring system; R⁴ is C₁₋₆alkyl; R⁵ is selected from hydrogen and C₁₋₆alkyl; R⁶ is selected from C₁₋₆alkyl, C₁₋₆alkyl(C₄₋₆cycloalkyl), C₁₋₆alkyl(C₁₋₆alkoxy), C₁₋₆alkylS(C₁₋₆alkyl), C₁₋₆alkylsulphonyl(C₁₋₆alkyl); (B) where q is an integer from 1 to 6 and R¹⁴ is halogeno; R⁷ is selected from C₁₋₆alkyl, C₁₋₆alkoxycarbonyl, C₂₋₆alkenyl, 1,3-benzodioxol-5-yl and aryl each optionally substituted by one or more substituents selected from C₁₋₆alkoxy, C₁₋₆alkyl, cyano, halogeno, and trifluoromethyl; R⁸ is aryl, heteroaryl, a bicyclic heteroaryl ring system linked to the nitrogen via a ring carbon or a 9 or 10 membered bicyclic ring system linked to the nitrogen via a ring carbon and each ring is optionally substituted with up to two substituents, which may be the same or different, and are selected from C₁₋₆alkyl, C₁₋₆alkoxy, C₁₋₆alkylthio, C₁₋₆alkyl(C₁₋₆alkoxy), C₁₋₆alkylaminoC₁₋₆alkyl, hydroxy, -CO₂H, -(CH₂)_pOH where p is 1 or 2, cyano, halogeno, and trifluoromethyl; R⁹ and R¹⁰ are each independently selected from hydrogen and C₁₋₆alkyl or R⁸ and R⁹ together with the nitrogen to which they are attached form a dihydroindolyl, or a dihydroquinolyl group; R¹¹ is selected from carboxyl, tetrazolyl, alkyl sulphonylcarbamyl, sulfo and sulfinyl; Y is oxygen, sulphur or sulfonyl; m is 0 or 1; and n is 0 or an integer from 1 to 4 with the proviso that when m and n cannot both be 0 and when m is 1, n is 0; or a pharmaceutically acceptable salt or <i>in vivo</i> hydrolysable ester thereof. The compounds inhibit the interaction of vascular cell-adhesion molecule-1 and fibronectin with integrin very late antigen 4 (α₄β₁). They have therapeutic applications such as in multiple sclerosis, rheumatoid arthritis, asthma, coronary artery disease and psoriasis.</p>		

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INTERNATIONAL SEARCH REPORT

Inter. Application No

PCT/GB 98/03334

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C07C275/34 C07D317/60 C07C317/50 C07D405/12 C07D407/12
C07D409/12 C07D417/12 C07D413/12 C07C323/60 C07K5/02
A61K31/195 A61K31/36 A61K31/41

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C07C C07D A61K C07K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 96 22966 A (BIOGEN INC) 1 August 1996 cited in the application see table 1; claims 1, 28-33 ---	1,6,8-10
A	WO 97 08145 A (G.D. SEARLE & CO) 6 March 1997 see page 4, line 18 - page 13, line 6; examples 108, 124 -----	1,6,8-10

☐ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

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- "&" document member of the same patent family

Date of the actual completion of the international search

12 May 1999

Date of mailing of the international search report

21/05/1999

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INTERNATIONAL SEARCH REPORT

International application No.

PCT/GB 98/03334

Box I Observations where certain claims were found unsearchable (Continuation of Item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.: 8-9
because they relate to subject matter not required to be searched by this Authority, namely:
Remark: Although claims 8-9
are directed to a method of treatment of the human/animal
body, the search has been carried out and based on the alleged
effects of the compound/composition.
2. ☐ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such
an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of Item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all
searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment
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covers only those claims for which fees were paid, specifically claims Nos.:
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restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

information on patent family members

International Application No

PCT/GB 98/03334

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
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